PATENT

Appl. No. 10/030,735

Amdt. dated November 11, 2005

Amendment and Reply under 37 CFR 1.116 Expedited

Procedure Examining Group 1644

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (Currently Amended) A peptide consisting of the sequence R₁-X₁-V R-X₂-R₂ R₁-X₁-X₂-X₃-X₄-R₂ or partial or full retro-inverso sequences thereof, wherein X₁ is selected from the group consisting of N, Q, and D; X₂ is V; X₃ is R; and X₄ is L; the X₁-V R-X₄ sequence is selected from the group consisting of N V R L (SEQ ID NO:57), N V R F (SEQ ID NO:51), Q V R L (SEQ ID NO: 80), Q V R F (SEQ ID NO:53), and D V R L (SEQ ID NO:102); R₁ is a hydrogen or from 1 to 6 amino acids, an acyl or an aryl group; and R₂ is from 1 to 3 amino acids, a hydroxide or an amide, provided that the peptide binds α3β1 integrin and does not comprise the sequence FQGVLQNVRFVF (SEQ ID NO:6).
- 2. (Currently amended) The peptide of claim 1, wherein the peptide contains the X_1 V R X_4 sequence X_1 - X_2 - X_3 - X_4 and is up to 12 amino acids in length.
- 3. (Previously presented) The peptide of claim 1 wherein R₁ is a peptide consisting of the sequence selected from the group consisting of FQGVLQ (SEQ ID NO:13), FAGVLQ (SEQ ID NO:14), FQGVAQ (SEQ ID NO:15), FQGVLA (SEQ ID NO:16), and FQGVLN (SEQ ID NO:17).
- 4. (Previously presented) A peptide that binds α3β1 integrin, wherein said peptide consists of a sequence selected from the group consisting of FQGVLQQVRFVF (SEQ ID NO:20), FQGVLQSVRFVF (SEQ ID NO:21), acQGVLQNVRF (SEQ ID NO:22), FQGVLNNVRFVF (SEQ ID NO:24), AQGVLQNVRFVF (SEQ ID NO:25), FAGVLQNVRFVF (SEQ ID NO:26), FQGVAQNVRFVF (SEQ ID NO:27),

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FQGVLQNVRFVA (SEQ ID NO:28), FQGVLANVRFVF (SEQ ID NO:29), FQGVLQNVRFV (SEQ ID NO:30), QGVLQNVRFVF (SEQ ID NO:31), and FQGVLQNVRF (SEQ ID NO:32).

5. (Currently amended) The A peptide of claim 1 consisting of the sequence R₁-X₁-X₂-X₃-X₄-R₂ or full retro-inverso sequences thereof, wherein X₁ is selected from the group consisting of N and Q; X₂ is V; X₃ is R; and X₄ is F; R₁ is a hydrogen or from 1 to 6 amino acids, an acyl or an aryl group; and R₂ is from 1 to 3 amino acids, a hydroxide or an amide, provided that the peptide binds α3β1 integrin, and wherein the X₁-V-R-X₄-X₁-X₂-X₃-X₄ portion of the sequence is optionally selected from the group consisting of NVRF (SEQ ID NO:51) and QVRF (SEQ ID NO:53).

6-7. (Cancel)

- 8. (Currently amended) A retro-inverso synthetic peptide consisting of the amino acid sequence, from C-terminal (left) to N-terminal (right): ri-R'₁-X'₁-X'₂-X'₃-X'₄-R'₂, wherein ri denotes a retro-inverso peptide sequence and all amino acids are D amino acids; wherein X₁ is selected from the group consisting of N, Q, and D; X₂ is V; X₃ is R; and X₄ is L; the X'₁-V-R-X'₄-sequence is selected from the group consisting of N V R-L (SEQ ID NO:57), N-V-R F (SEQ ID NO:51), Q-V-R-L (SEQ ID NO: 80), Q-V-R F (SEQ ID NO:53), and D-V-R-L (SEQ ID NO:102); R'₁ is a hydrogen or from 1 to 6 amino acids, a hydroxide or an amide; and R'₂ is from 1 to 3 amino acids, an acyl or an aryl group.
- 9. (Currently amended) The peptide of claim 8, wherein the peptide contains the $\frac{X_1^2-X_2^2-X_3^2-X_4^2}{X_1^2-X_2^2-X_3^2-X_4^2}$ and is up to 12 amino acids in length.
- 10. (Previously presented) A peptide consisting of the sequence FQGVLQNVRFVF (SEQ ID NO:6) wherein every amino acid in said sequence is a D-amino acid.

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10. (Previously presented) A peptide consisting of the sequence FQGVLQNVRFVF (SEQ ID NO:6) wherein every amino acid in said sequence is a D-amino acid.

11-12. (Canceled)

- 13. (Previously presented) A composition comprising a peptide according to claim 1 and a pharmaceutically acceptable carrier.
- 14. (Previously presented) A composition comprising a peptide according to claim 1 in a sterile aqueous solution.

15-19. (Canceled)

- 20. (Withdrawn) An in vitro method of inhibiting adhesion of a cell expressing $\alpha 3\beta 1$ integrin to an extracellular matrix comprising contacting said cell with a peptide according to claim 1.
- 21. (Withdrawn) The method of claim 20 wherein the extracellular matrix comprises TSP1 or laminins.

22. (Cancel)

- 23. (Withdrawn) The method of claim 20 wherein said cell comprises an epithelial or an endothelial cell.
 - 24. (Withdrawn) The method of claim 20 wherein said cell is a tumor cell.

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- 25. (Withdrawn) The method of claim 20 wherein said cell is a breast carcinoma cell or a small cell lung carcinoma.
- 26. (Withdrawn) An in vitro method of inhibiting 03\$1 integrin-mediated cell motility, comprising contacting a cell with a peptide according to claim 1.
 - 27. (Canceled)
- 28. (Withdrawn) The method of claim 26 wherein the cell is an epithelial cell, an endothelial cell or a malignant cell.
- 29. (Withdrawn) An *in vitro* method of inhibiting proliferation of endothelial cells, comprising contacting said cells with a peptide according to claim 1.
- 30. (Withdrawn) An *in vitro* method of inhibiting proliferation of small cell lung carcinoma cells, comprising contacting said cells with a peptide according to claim 2.

31-45. (Canceled)

- 46. (Currently amended) A peptide consisting of the sequence R_1 -D-V-R-F- R_{27} R_1 - X_1 - X_2 - X_3 - X_4 - R_2 or partial or full retro-inverso sequences thereof, wherein D-V-R-F is SEQ ID NO:54; X_1 is D; X_2 is V; X_3 is R; and X_4 is F; R_1 is a hydrogen or from 1 to 6 amino acids, an acyl or an aryl group; and R_2 is 2 or 3 amino acids, a hydroxide or an amide, provided that the peptide binds $\alpha 3\beta 1$ integrin.
- 47. (Previously presented) The peptide according to claim 46 consisting of the sequence FQGVLQDVRFVF (SEQ ID NO:19).

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- 48. (Previously presented) The peptide of claim 46, wherein the peptide contains the sequence DVRF (SEQ ID NO:54) and is up to 12 amino acids in length.
- 49. (Previously presented) The peptide of claim 46 wherein R₁ is a peptide consisting of the sequence selected from the group consisting of FQGVLQ (SEQ ID NO:13), FAGVLQ (SEQ ID NO:14), FQGVAQ (SEQ ID NO:15), FQGVLA (SEQ ID NO:16), and FQGVLN (SEQ ID NO:17).
- 50. (Previously presented) The peptide of claim 46 that contains at least one D-amino acid.
- 51. (Previously presented) A composition comprising a peptide according to claim 46 and a pharmaceutically acceptable carrier.
- 52. (Previously presented) A composition comprising a peptide according to claim 46 in a sterile aqueous solution.
- 53. (Previously presented) A retro-inverso synthetic peptide consisting of the amino acid sequence, from C-terminal (left) to N-terminal (right): ri- R'₁-D-V-R-F-R'₂, wherein ri denotes a retro-inverso peptide sequence and all amino acids are D amino acids and D-V-R-F is SEQ ID NO:54; R'₁ is a hydrogen or from 1 to 6 amino acids, a hydroxide or an amide; and R'₂ is 2 or 3 amino acids, a hydroxide or an amide, provided that the peptide binds α 3 β 1 integrin.
- 54. (Previously presented) The peptide of claim 46, wherein the peptide contains the sequence DVRF (SEQ ID NO:54) and is up to 12 amino acids in length.